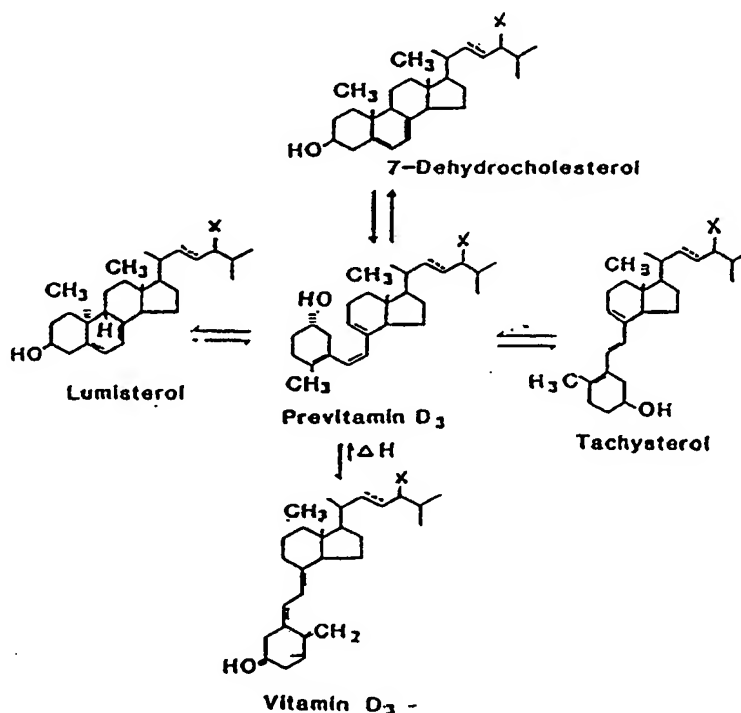




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(54) Title: COMPOSITIONS COMPRISING VITAMIN D PRECURSORS, ANALOGS THEREOF AND THEIR USE**(57) Abstract**

Methods for providing active-type vitamin D compounds to an individual are disclosed. The individuals are exposed to sunlight to produce vitamin D and analogs or derivatives thereof via the skin. Pharmaceutical compositions comprising lumisterol and/or tachysterol and analogs or derivatives thereof are also disclosed.

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TITLE OF THE INVENTION

COMPOSITIONS COMPRISING VITAMIN D PRECURSORS,
ANALOGS THEREOF AND THEIR USEField of the Invention

5 The invention is in the field of cosmetics and
medicinal chemistry. In particular, the present
invention relates to topical compositions which
provide vitamin D, derivatives and analogs thereof
throughout the year. In another aspect, the present
10 invention relates to a method of producing previtamin
D, derivatives and analogs thereof. The topical
compositions of the invention allow a user in the high
northern and southern latitudes to produce previtamin
D, derivatives and analogs thereof in and on their
15 skin even when exposed to low energy sunlight in the
winter as well as in the morning and evening
throughout the year. The method employs tachysterol
and lumisterol, derivatives and analogs thereof which
photoisomerize to previtamin D, derivatives and
20 analogs thereof, respectively, when exposed to low
levels of ultraviolet radiation.

Background of the Invention

25 Vitamin D₃ is a derivative of provitamin D₃ (7-
dehydrocholesterol), the immediate biological
precursor of cholesterol. With adequate exposure to
sunlight, dietary supplements are not normally
required. Holick et al. in Braunwald et al.,
Harrison's Principles of Internal Medicine, 11th ed.
McGraw-Hill (1987), pp. 1857-69. However, not all

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individuals are exposed to the adequate levels of sunlight, especially in the winter.

When skin is exposed to sunlight or artificial sources of ultraviolet (UV) radiation, the UV radiation penetrates the epidermis and causes a variety of biochemical reactions. Included in these reactions are the transformation of provitamin D₃ to vitamin D₃. The electromagnetic energy having wavelengths between 290 and 315 nm is absorbed by provitamin D₃ resulting in its fragmentation to previtamin D₃. Although previtamin D₃ is biologically inert, it is thermally labile and spontaneously undergoes a temperature-dependent rearrangement to form the thermally stable vitamin D₃. After biosynthesis, vitamin D₃ is translocated from the epidermis into the circulation via a vitamin-D binding protein. Holick et al., Science 211:590-593 (1981); Holick et al. in Braunwald et al., Harrison's Principles of Internal Medicine, 11th ed., McGraw-Hill (1987), pp. 1857-69.

Factors that are frequently considered as affecting the cutaneous synthesis of vitamin D₃ include age, altitude, geographical location, time of day and area of exposure to sunlight. Common to most of these factors is the availability of the requisite amount of ultraviolet radiation with energies between 290 and 315 nm which is necessary to convert provitamin D₃ to vitamin D₃. MacLaughlin et al., Science 216:1001-1003 (1982).

The availability of vitamin D precursor in the skin and its photo-induced transformation to previtamin D₃ and then to vitamin D₃ is an efficient

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physiological source of and mechanism for the replenishment of vitamin D₃. However, during the winter in northern latitudes, sunlight does not contain enough high energy ultraviolet radiation to convert provitamin D₃ (7-dehydrocholesterol) in human skin to previtamin D₃ (Webb, Kline and Holick, J. Clin. Endocrinol Met. 57:373-378 (1988)). As a result, individuals in these latitudes cannot make vitamin D₃ in their skin, even when they are exposed to sunlight. The lack of adequate exposure to ultraviolet radiation gives rise to the possibility of serious vitamin D deficiency, a breakdown in blood calcium regulation with concomitant hypocalcemia and bone calcium wasting.

The availability of the vitamin D precursor in the skin and its photo-induced transformation to previtamin D₃, and then to vitamin D₃, is an efficient physiological source of, and mechanism for the replenishment of vitamin D₃. Previously, it was thought that the only method of producing previtamin D₃ was to transform provitamin D₃. This transformation requires sunlight or artificial UV light in the region of 290-315 nm. Therefore, in areas where the available light energy is below this range (wavelengths greater than 316 nm), the transformation does not occur to any significant extent. Kobayashi et al., J. Nutr. Sci. Vitaminol. 19:123 (1973).

It has been disclosed (Holick, M., Transactions of the Association of American Physicians, 42:54-63 (1979); Molecular Endocrinology; MacIntyre and Szelke, eds.; Elsevier/North Holland Biomedical Press (1979), pp.301-308) that the topical application of

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hydroxylated metabolites of provitamin D compounds to the skin combined with U.V. phototherapy is a method for the sustained administration of vitamin D metabolites to patients who suffer vitamin D metabolic disorders. When the hydroxylated provitamins are applied and irradiated with ultraviolet radiation, they convert to hydroxylated previtamins which then thermally isomerize to the hydroxylated vitamin D. This work is also disclosed in Holick et al., New England Journal of Medicine 301:349-354 (1980) and U.S. Patent No. 4,310,511 (Jan. 12, 1982).

1,25-Dihydroxyvitamin D₃ and its analogs have been shown to be powerful antiproliferative agents which are effective for the treatment of the hyperproliferative disorder psoriasis (DeLuca, H. Fed. Proc. Am. Soc. Biol. 2:224-236 (1988); Holick in DeGroot et al., Endocrinology 2:902-926, Grune and Stratton, N.Y., N.Y., (1988); Morimoto et al., Br. J. Dermatol. 115:421-429 (1986); Holick, Arch. Dermatol. 125:1692-1697 (1989)).

Hungarian Patent No. 102,939 discloses cosmetic creams containing provitamin D (such as ergosterol) which, when irradiated with ultraviolet rays, are transformed into vitamin D.

MacLaughlin et al., Science 216:1001-1003 (1982), disclose the synthesis of previtamin D₃ from provitamin D₃ in human skin and in an organic solvent after exposure to narrow-band radiation or simulated solar radiation. When human skin or an organic solvent containing provitamin D₃ were exposed to 295 nm radiation, up to 65% of the provitamin D₃ was converted to previtamin D₃. The authors further disclose that

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the optimum wavelength for the production of previtamin D₃ is between 295 nm and 300 nm.

Dauben et al., J. Am. Chem. Soc. 104:5780-5781 (1982); J. Am. Chem. Soc. 104:355-356 (1982), disclose the effect of wavelength on the photochemistry of provitamin D₃ and the effect of wavelength on the production of previtamin D₃. The authors found that when provitamin D₃ is exposed to light in the range of 254 nm, it is converted to a variety of photoproducts, the major portion being about 75% tachysterol. This mixture was then exposed to either 300 nm of light, broad-band 350 nm light or 355 nm light to give a build up of previtamin D₃. Dauben et al. conclude that if provitamin D₃ is first irradiated at 0°C with 254 nm light to give a quasi photostationary state of provitamin D₃, previtamin D₃, tachysterol and lumisterol, and the mixture is thereafter irradiated (0°C) with 350 nm light, a maximum of 83% previtamin D₃ is produced.

Malatesta et al., J. Amer. Chem. Soc. 103:6781-6783 (1981), disclose the effects of different UV wavelengths on the relative quantities of photoproducts produced from provitamin D₃.

Holick et al. disclose that the photochemical conversion of previtamin D₃ to lumisterol and tachysterol is the major factor that prevents vitamin D₃ intoxication after a single prolonged exposure to the sun. Holick et al., Science 211:590-592 (1981). The corollary to this finding is that lumisterol and tachysterol are two biologically inert products thought to be sloughed off the skin during the natural turnover of the epidermal cells.

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Provitamin D₂ (ergosterol) is the precursor of vitamin D₂. Vitamin D₂ is one of the major forms of vitamin D that is used to fortify foods such as milk and multivitamins.

5

SUMMARY OF THE INVENTION

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The present invention is related to the discovery that topical formulations comprising lumisterol and tachysterol, analogs and derivatives thereof are effective means of providing previtamin D, derivatives and analogs thereof to individuals. The present invention utilizes the low energy UV photoconversion of lumisterol and tachysterol, analogs or derivatives thereof, to previtamin D, analogs or derivatives thereof, respectfully, as a method of producing vitamin D, analogs or derivatives thereof in the skin. It is this novel finding that solves the problem of producing vitamin D compounds via the skin in areas of low energy sunlight.

15

20

In particular, the invention is directed to lumisterol and tachysterol, derivatives and analogs thereof which are convertible to vitamin D analogs in the presence of low energy UV light. The invention is also directed to pharmaceutical compositions containing an effective amount of lumisterol and/or tachysterol, derivatives or analogs thereof, and a pharmaceutically effective carrier.

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The invention is also directed to a method for providing vitamin D, analogs or derivatives thereof, to an individual by administering to the individual a pharmaceutical composition of the invention.

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The invention is also directed to a method of treating hyperproliferative disorders of the skin including psoriasis, healing wounds and inhibiting scar formation with the pharmaceutical compositions of the invention.

The invention is also directed to the treatment of ulcers such as diabetic ulcers of the foot, decubitus ulcers (bed sores), genito-urinary ulcers, and ulcerative keratitis with the pharmaceutical compositions of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 illustrates the photochemical conversion of provitamin D to vitamin D and the concomitant production of lumisterol and tachysterol. When the bond between C-22 and C-23 is a single covalent bond and X is hydrogen, the compounds belong to the D₃ family, e.g. vitamin D₃. Where the bond between C-22 and C-23 is a double covalent bond and X is methyl, the compounds belong to the D₂ family, e.g. vitamin D₂.

Figure 2 depicts an HPLC trace of a control solution of provitamin D₃ (A) and a solution of provitamin D₃ exposed to sunlight on a day during the winter (B).

Figure 3 depicts an HPLC trace of a control solution of tachysterol (A) and a solution of tachysterol exposed to sunlight on a day during the winter (B).

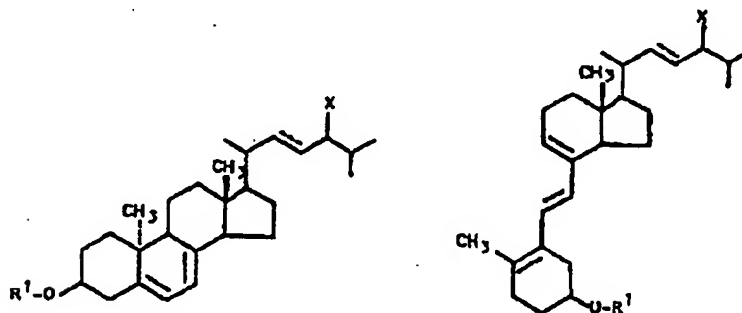
Figure 4 depicts an HPLC trace of a control solution of lumisterol (A) and a solution of

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lumisterol exposed to sunlight on a day during the winter (B).

DESCRIPTION OF THE PREFERRED EMBODIMENTS

5 The active compounds utilized in the present invention are tachysterol, lumisterol and derivatives thereof, either alone or in combination. The tachysterol and lumisterol derivatives have the following Formulae (I) and (II), respectively:



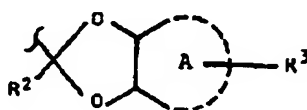
(I)

(II)

10 wherein the bond between C-22 and C-23 is a single or double bond;
X is hydrogen, methyl or ethyl; and

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R¹ is hydrogen or a straight or branched chain glycosidic residue containing 1-20 glycosidic units per residue, or R¹ is an orthoester glycoside moiety of the Formula (III):



5

(III)

where A represents a glucofuranosyl or a glucopyranosyl ring;

R² is hydrogen, lower (C₁-C₄) alkyl, C₇-C₁₀ aralkyl, or C₆-C₁₀ aryl; and

10

R₃ is hydrogen or a straight or branch chain glycosidic residue containing 1-20 glycosidic units per residue.

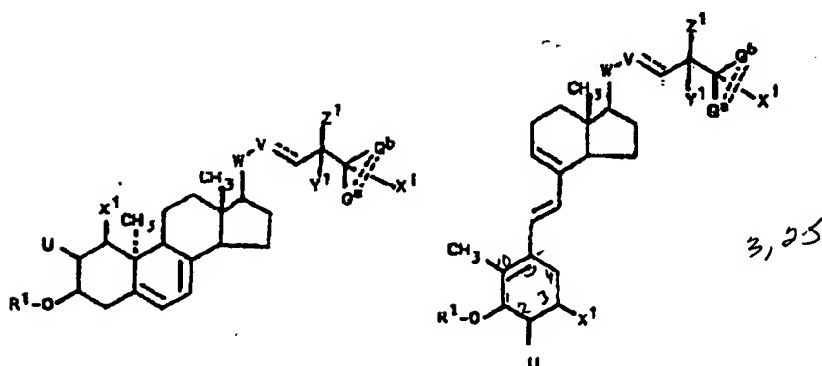
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These compounds are photoisomers of previtamin D, the precursor of biologically active vitamin D. Tachysterol and lumisterol may be prepared by photoisomerization and isolation as disclosed by Holick et al., Biochem. 18:1003-1008 (1979). Analogous methods for making the corresponding glycosidic and orthoester glycoside derivatives are taught, for example, by Holick et al., U.S. Patent Nos. 4,410,515 and 4,521,410, the disclosures of which are fully incorporated by reference herein.

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The tachysterol and lumisterol analogs of the present invention have the following Formulae (IV) and (V), respectively:



(IV)

(V)

5

wherein the bond between carbons C-22 and C-23 is single or double bond;

Y^1 is hydrogen, F, CH_3 , CH_2CH_3 or X^1 ;

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U is hydrogen, -OH or -O-(C₂-C₄ alkyl)-OH;

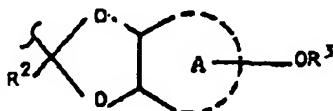
Z¹ is F, H or X¹;

Q^a is CF₃ or CH₂X¹;

Q^b is CF₃ or CH₃;

wherein X¹ is selected from the group consisting of hydrogen, -OH and OR¹;

wherein R¹ is hydrogen or a straight or branched chain glycosidic residue containing 1-20 glycosidic units per residue, or R¹ is an orthoester glycoside moiety of the Formula (III):



(III)

wherein A represents a glucofuranosyl or glucopyranosyl ring;

R² is hydrogen, lower C₁-C₄ alkyl or aryl, with the proviso that aryl is phenyl or phenyl substituted by chloro, fluoro, bromo, iodo, lower C₁-C₄ alkyl, C₁-C₄ alkoxy; or naphthyl; and

R³ is hydrogen or a straight or branched chain glycosidic residue containing 1-20 glycosidic units per residue;

W is CH-CH₃ or O; -

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V is CH₂ or O;
with the proviso that both W and V are not both O; and
"==" is either a single bond between Q^a and Q^b or
a hydrogen atom on Q^a and Q^b.

5 These compounds are photoisomers of previtamin D
analogs, the precursor of biologically active vitamin
D analogs.

10 Examples of particular vitamin D analogs are
taught, for example, by Holick et al., U.S. Patent No.
4,310,511 (Jan. 12, 1982); Partridge et al., U.S.
Patent No. 4,634,692 (1987); Yamada, JP Publication
No. J5 5111-460; DeLuca et al., U.S. Patent No.
4,719,205 (1988); Holick et al., U.S. Patent No.
15 4,410,515 (1983); Holick et al., U.S. Patent No.
4,521,410 (1985); Holick et al., U.S. Patent No.
4,230,701; and Shiina et al., Arch. Biochem. Biophys.
220:90 (1983), the disclosures of which are fully
incorporated by reference herein. Methods for making
the corresponding glycosidic and orthoester glycoside
20 vitamin D analogs are taught, for example, by Holick
et al., U.S. Patent Nos. 4,410,515 and 4,521,410, the
disclosures of which are fully incorporated by
reference herein.

25 The tachysterol and lumisterol analogs may be
prepared by photoisomerization of the requisite
provitamin D analog according to the method disclosed
by Holick et al., Biochem. 18:1003-1008 (1979).

30 By administering an effective amount of
tachysterol, lumisterol and analogs or derivatives
thereof in topical compositions according to this
invention, it is possible for the first time to
provide a method which allows individuals living in

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regions of low energy sunlight to produce vitamin D compounds via their skin, thus preventing harmful vitamin D₃ depletion. The compositions of the present invention may be used, therefore, in methods of treating or preventing osteomalacia due to vitamin D deficiency, and calcium disorders resulting from a lack of vitamin D (a lack of vitamin D leads to deficient intestinal absorption of calcium which results in hypocalcemia), glucocorticoid-induced decrease in calcium absorption, osteoporosis, senile decrease in calcium absorption, hypoparathyroidism, milk fever disease, turkey weak leg disease, etc.

The present invention also provides for a method of healing wounds and inhibiting scar formation and treating hyperproliferative disorders of the skin including psoriasis by administering an effective amount of a tachysterol or lumisterol analog of the invention. Wounds to the external epithelium include cuts, punctures and lacerations, including corneal lacerations. The invention also provides for the treatment of ulcers such as diabetic ulcers of the feet, decubitus ulcers (bed sores), genito-urinary ulcers, and ulcerative keratitis by administering an effective amount of a tachysterol or lumisterol analog of the invention. Ulcerative keratitis is caused, for example, by extended wear of contact lenses.

Genito-urinary ulcers treatable with the tachysterol and lumisterol analogs of the invention include those caused by, for example, herpes simplex virus as well as other viral, fungal and bacterial infections. See Harrison's Principles of Internal

Medicine, E. Braunwald et al. (eds.); McGraw-Hill Book Co., New York, N.Y., 1987, pp. 514-516.

Examples of tachysterol and lumisterol analogs include 1-hydroxytachysterol₂, 1-hydroxytachysterol₃, 1-hydroxylumisterol₂, 1-hydroxylumisterol₃, 1,24-dihydroxytachysterol₂, 1,24-dihydroxytachysterol₃, 1,24-dihydroxylumisterol₂, 1,24-dihydroxylumisterol₃, 1,25-dihydroxytachysterol₂, 1,25-dihydroxytachysterol₃, 1,25-dihydroxylumisterol₂, 1,25-dihydroxylumisterol₃, 24,25-dihydroxytachysterol₂, 24,25-dihydroxytachysterol₃, 24,25-dihydroxylumisterol₂, 24,25-dihydroxylumisterol₃, 25,26-dihydroxytachysterol₂, 25,26-dihydroxytachysterol₃, 25,26-dihydroxylumisterol₂, 25,26-dihydroxylumisterol₃, 1,24,25-trihydroxytachysterol₂, 1,24,25-trihydroxytachysterol₃, 1,24,25-trihydroxylumisterol₂, 1,24,25-trihydroxylumisterol₃, 2-β-(3-hydroxypropoxy)-1 α,25-dihydroxytachysterol₂, 2-β-(3-hydroxypropoxy)-1 α,25-dihydroxytachysterol₃, 2-β-(3-hydroxypropoxy)-1 α,25-dihydroxylumisterol₂, 2-β-(3-hydroxypropoxy)-1 α,25-dihydroxylumisterol₃, as well as the side chain fluoro derivatives of 1,25-dihydroxytachysterol₂, 1,25-dihydroxytachysterol₃, 1,25-dihydroxylumisterol₂, 1,25-dihydroxylumisterol₃, 1-hydroxytachysterol₂, 1-hydroxytachysterol₃, 1-hydroxylumisterol₂, and 1-hydroxylumisterol₃. Also included are the 20- and 22-oxa tachysterol and lumisterol derivatives including 20-oxa-1α(OH)tachysterol₂, 20-oxa-1α(OH)tachysterol₃, 20-oxa-1α(OH)lumisterol₂, 20-oxa-1α(OH)lumisterol₃, 20-oxa-1α,25(OH)₂tachysterol₂, 20-oxa-1α,25(OH)₂tachysterol₃, 20-oxa-1α,25(OH)₂lumisterol₂, 20-oxa-1α,25(OH)₂-

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lumisterol₃, 22-oxa-1 α (OH)tachysterol₂, 22-oxa-1 α (OH)tachysterol₃, 22-oxa-1 α (OH)lumisterol₂, 22-oxa-1 α (OH)lumisterol₃, 22-oxa-1 α ,25(OH)₂tachysterol₂, 22-oxa-1 α ,25(OH)₂tachysterol₃, 22-oxa-1 α ,25(OH)₂lumisterol₂ and 22-oxa-1 α ,25(OH)₂lumisterol₃. Also included within the scope of the present invention are 25,26 cyclopropyl compounds including 1,24-dihydroxy-25,26-dehydrotachysterol₃, 1,24-dihydroxy-25,26-dehydrotachysterol₂, 1,24-dihydroxy-25,26-dehydrolumisterol₃, and 1,24-dihydroxy-25,26-dehydrolumisterol₂.

Foremost among the individuals which may be treated with the compositions of the invention are humans, although the invention is not intended to be so limited. Any animal which may benefit from treatment with the compositions of the invention are within the spirit and scope of the present invention.

By using tachysterol and lumisterol analogs in topical compositions according to this invention, it is possible for the first time to provide a method which allows individuals living in regions of low sunlight to produce vitamin D analogs via their skin. The compositions of the present invention may be used, therefore, in methods of treating decubitus and diabetic foot ulcers; ulcerative keratitis; treating psoriasis; wound healing; inhibiting scar formation; treating or preventing osteodystrophy due to an acquired or inherited disorder in vitamin D metabolism; glucocorticoid-induced decrease in calcium absorption; osteoporosis; senile decrease in calcium absorption; hypoparathyroidism; milk fever disease; and turkey weak leg disease.

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The compounds of the present invention can be administered in any appropriate pharmacological carrier for topical or intravenous administration. The dosage administered will be dependent on the age, health and weight of the recipient, and the nature of the effect desired.

The topical compositions of the invention may be applied so that at least 0.1 microgram, preferably at least about 10 micrograms to about 100 mg of the vitamin D precursor/gm carrier is administered to the skin. A preferred range is between about 1 microgram to about 1 milligram of tachysterol, lumisterol or analog or derivative thereof/gm carrier.

The compositions of the invention formulated for intravenous administration may comprise at least about 0.1 microgram, preferably at least about 1.0 microgram to about 100 mg of the vitamin D precursor or analog precursor per ml of physiologically acceptable solution. A most preferred range is about 1.0 micrograms to about 100 micrograms of tachysterol, lumisterol or analog or derivative thereof per ml of solution.

The compounds can be employed in a pharmacologically inert topical carrier such as one comprising a gel, an ointment or a cream, including such carriers as water, glycerol, alcohol, propylene glycol, fatty alcohols, triglycerides, fatty acid esters or mineral oils. Other possible carriers are liquid petrolatum, isopropylpalmitate, polyethylene glycol ethanol 95%, polyoxyethylene monolaurate 5% in water, sodium lauryl sulfate 5% in water, and the like. Minerals such as

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anti-oxidants, humectants, viscosity stabilizers and the like may be added, if necessary.

Alternatively, the compounds may be employed as part of a sun screen lotion which selectively screens the harmful high energy UV radiation (below 315 nm) but which allows medium and low energy UV radiation (above 315 nm) to pass which is of sufficient energy to photoisomerize lumisterol, tachysterol and the analogs or derivatives thereof to previtamin D or the analogs or derivatives thereof. Alternatively, the compounds of the invention may be added to broad range sun screens that absorb radiation with energies of up to 360 nm. Such sun screen lotions may comprise any of those well known to those of ordinary skill in the art, for example, ethyl p-aminobenzoate (benzocaine), p-aminobenzoic acid (PABA), octyl methoxycinnamate (PARASOL[®] MCX), butyl methoxydibenzoylmethane (PARASOL[®] 1789), phenyl salicylate (salcol), 2-ethoxyethyl p-methoxycinnamate, glyceryl p-aminobenzoate, 2,4-dibenzoyl resorcinol, octyl dimethyl PABA, oxybenzone, benzophenones, methyl anthranilate, cinoxate, amyldimethyl PABA, homomenthyl salicylate, digalloyl trioleate, ethyl-p-glycosylimido benzoate, and red veterinary petrolatum. For other examples, see Algra et al., Int. J. Derm. 17:628-634 (1978), Sayre, R.M. et al., Photochem. Photobiol. 29:559-566 (1979).

Preparations for parenteral administration include sterile or aqueous or non-aqueous solutions, suspensions, and emulsions. Examples of non-aqueous solvents are propylene glycol, polyethylene glycol, vegetable oils such as olive oil, and injectable organic esters such as ethyl oleate. Aqueous carriers

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include water, alcoholic/aqueous solutions, emulsions or suspensions, including saline and buffered media. Parenteral vehicles include sodium chloride solution, Ringer's dextrose, dextrose and sodium chloride, lactated Ringer's, or fixed oils. Intravenous vehicles include fluid and nutrient replenishers, electrolyte replenishers, such as those based on Ringer's dextrose, and the like. Preservatives and other additives may also be present, such as, for example, antimicrobials, anti-oxidants, chelating agents, inert gases and the like. See, generally, Remington's Pharmaceutical Science, 16th Ed., Mack Eds., 1980.

The compositions comprising tachysterol and/or lumisterol and analogs or derivatives thereof which are formulated for parenteral administration may be utilized to provide an individual with these vitamin D analog precursors so as to allow the production of vitamin D analogs or derivatives in the skin in the presence of medium and low energy UV radiation.

The invention further relates to solutions comprising the tachysterol and lumisterol, analogs and derivatives thereof which may be exposed to UV radiation to allow the preparation of a solution comprising an active vitamin D compound as desired just before administration to the individual. This method avoids the decomposition of vitamin D and analogs thereof which occurs in solution. Solutions which may comprise compounds of the invention may include the above-listed parenteral solutions. Of course, the solutions comprising the lumisterol and tachysterol analogs must be stored in an opaque

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container to avoid premature conversion of tachysterol and lumisterol analogs to the corresponding vitamin D analog.

Having now generally described this invention, the same will be understood by reference to an example which is provided herein for purposes of illustration only and is not intending to be limited unless otherwise specified.

Example 1

Crystalline provitamin D₃ was dissolved in methanol at a concentration of 10 micrograms/ml. Ten ml of this solution was placed in quartz test tubes. One test tube containing provitamin D₃ in methanol was exposed to direct sunlight in Boston during November, 1989 between 9 AM and 10 AM (Fig. 2B) while a similar sample remained in the dark over the same period of time (Fig. 2A). At the end of the exposure, a small aliquot was taken from each test tube and chromatographed on a high performance liquid chromatograph according to MacLaughlin et al., Science 216:1001-1003 (1982). Similar studies were conducted with lumisterol (Fig. 4) and tachysterol (Fig. 3) that were prepared as previously described (Holick et al., Biochem. 18:1003-1008 (1979)). The analysis of all the chromatograms in Figs. 2-4 revealed that when tachysterol and lumisterol were exposed to sunlight in November between 9 and 10 AM, they underwent photoisomerization to previtamin D₃ (Figs. 3B, 4B). In contrast, provitamin D₃ exposed to the same direct sunlight did not convert to previtamin D₃ (Fig. 2B). All samples that were kept in the dark for the same

-20-

time did not convert to previtamin D₃ (Figs. 2A, 3A, 4A).

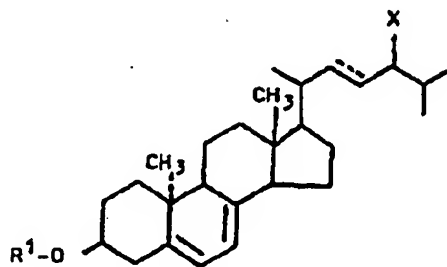
5 It is expected that the tachysterol and lumisterol analogs of the present invention, upon irradiation with the same low intensity and energy UV light, will give the corresponding analogs.

10 Having now generally described this invention, it will be apparent to one of ordinary skill in the art that the same can be carried out in a variety of embodiments and variations which are equivalent without affecting the spirit or scope of the invention or any embodiments thereof.

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WHAT IS CLAIMED IS:

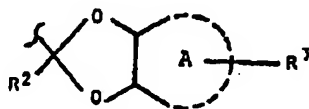
1. A composition comprising a pharmaceutically acceptable carrier and a compound of the formula



wherein the bond between C-22 and C-23 is a single or double bond;

X is hydrogen, methyl or ethyl; and

R¹ is hydrogen or a straight or branched chain glycosidic residue containing 1-20 glycosidic units per residue, or R¹ is an orthoester glycoside moiety of the formula:



where A represents a glucofuranosyl or a glucopyranosyl ring;

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R² is hydrogen, lower (C₁-C₄) alkyl, C₇-C₁₀ aralkyl, or C₆-C₁₀ aryl; and

R₃ is hydrogen or a straight or branch chain glycosidic residue containing 1-20 glycosidic units per residue;

wherein said compound is present in an amount effective to provide vitamin D₃ when said composition is administered to an individual.

2. The composition of claim 1, wherein said compound is lumisterol₃.

3. The composition of claim 1, wherein said carrier is effective for topical administration.

4. The composition of claim 3, further comprising one or more sun screen agents.

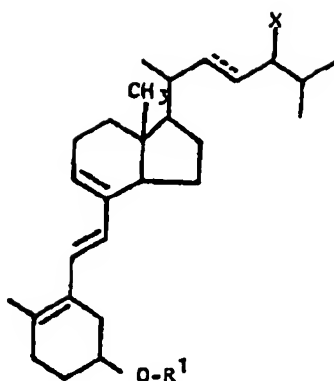
5. The composition of claim 1, wherein said carrier is effective for parenteral administration.

6. The composition of claim 1, wherein said compound is present in an amount of 0.00001 to 10% by weight.

7. The composition of claim 1, wherein said compound is present in an amount of 0.0001 to 0.01% by weight.

8. A composition a pharmaceutically acceptable carrier and a compound having the formula:

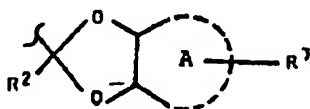
-23-



wherein the bond between C-22 and C-23 is a single or double bond;

X is hydrogen, methyl or ethyl; and

R¹ is hydrogen or a straight or branched chain glycosidic residue containing 1-20 glycosidic units per residue, or R¹ is an orthoester glycoside moiety of the formula:



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where A represents a glucofuranosyl or a glucopyranosyl ring;

R² is hydrogen, lower (C₁-C₄) alkyl, C₇-C₁₀ aralkyl, or C₆-C₁₀ aryl; and

5 R₃ is hydrogen or a straight or branch chain glycosidic residue containing 1-20 glycosidic units per residue;

10 wherein said compound is present in an amount effective to provide vitamin D₃ when said composition is administered to an individual.

9. The composition of claim 8, wherein said compound is tachysterol₃.

10. The composition of claim 8, wherein said carrier is effective for topical administration.

15 11. The composition of claim 10, further comprising one or more sun screen agents.

12. The composition of claim 8, wherein said carrier is effective for parenteral administration.

20 13. The composition of claim 8, wherein said compound is present in an amount of 0.00001 to 10% by weight.

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14. The composition of claim 8, wherein said compound is present in an amount of 0.0001 to 0.01% by weight.

5 15. A composition comprising lumisterol and tachysterol and a pharmaceutically acceptable carrier wherein said lumisterol and tachysterol are present in an amount effective to provide vitamin D₃ when said composition is administered to an individual.

10 16. The composition of claim 15, wherein said carrier is effective for topical administration.

17. The composition of claim 15, wherein said carrier is effective for parenteral administration.

15 18. The composition of claim 15, wherein said lumisterol and tachysterol are individually present in an amount of from 0.00001 to 10% by weight.

19. The composition of claim 15, wherein said lumisterol and tachysterol are individually present in an amount of from 0.001 to 0.01% by weight.

20 20. A method for providing vitamin D₃ to an individual which comprises administering to said individual the pharmaceutical composition of any one of claims 1, 8 and 15 and exposing said individual to UV radiation.

25 21. The method of claim 20, wherein said composition is administered by topical means.

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22. The method of claim 21, wherein said composition further comprises one or more sun screen agents.

5 23. The method of claim 20, wherein said composition is administered by intravenous means.

24. The method of claim 20, wherein said UV radiation is provided by sunlight of insufficient intensity and wavelength to effect the conversion of provitamin D to vitamin D.

10 25. The method of claim 24, wherein said UV radiation has a wavelength above 315 nm.

15 26. A method for treating or preventing osteomalacia due to vitamin D deficiency or a calcium disorder resulting from a lack of vitamin D, glucocorticoid-induced decrease in calcium absorption, osteoporosis, senile decrease in calcium absorption, hypoparathyroidism, milk fever disease, or turkey weak leg disease in an individual which comprises administering to said individual the pharmaceutical
20 composition of any one of claims 1, 8 and 15 and exposing said individual to low energy UV radiation.

27. The method of claim 26, wherein said composition is administered by topical means.

25 28. The method of claim 27, wherein said composition further comprises one or more sun screen agents.

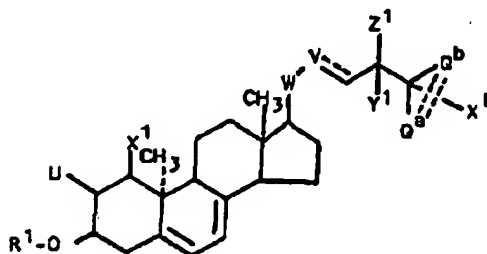
-27-

29. The method of claim 26, wherein said composition is administered by intravenous means.

30. The method of claim 26, wherein said UV radiation is provided by sunlight of insufficient intensity and wave-length to effect the conversion of provitamin D to vitamin D.

31. The method of claim 30, wherein said UV radiation has a wavelength above 315 nm.

32. A composition comprising a pharmaceutically acceptable carrier and a compound of the formula



wherein the bond between carbons C-22 and C-23 is single or double bond;

Y^1 is hydrogen, F, CH_3 , CH_2CH_3 or X^1 ;

U is hydrogen, -OH or -O-(C₂-C₄ alkyl)-OH;

z^1 is F, H or x^1 ;

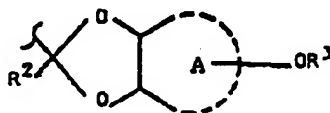
Q^a is CF_3 or CH_2X^1 ;

Q^b is CF_3 or CH_3 ;

wherein X' is selected from the group consisting of hydrogen, -OH and OR¹;

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wherein R^1 is hydrogen or a straight or branched chain glycosidic residue containing 1-20 glycosidic units per residue, or R^1 is an orthoester glycoside moiety of the formula:



wherein A represents a glucofuranosyl or glucopyranosyl ring;

R^2 is hydrogen, lower alkyl, or aryl, with the proviso that aryl is phenyl or phenyl substituted by chloro, fluoro, bromo, iodo, lower C_1 - C_4 alkyl, C_1 - C_4 alkoxy; or naphthyl; and

R^3 is hydrogen or a straight or branched chain glycosidic residue containing 1-20 glycosidic units per residue;

W is $CH-CH_2$ or O;

V is CH_2 or O;

with the proviso that both W and V are not both O; and

"=" is either a single bond between Q^a and Q^b or a hydrogen atom on Q^a and Q^b ;

wherein said compound is present in an amount effective to provide a vitamin D analog when said composition is administered to an individual.

33. The composition of claim 32, wherein said compound is $1\alpha,25$ -dihydroxylumisterol₃, $1\alpha,25$ -dihydroxylumisterol₂, 1α -hydroxylumisterol₃, 1α -hydroxylumisterol₂, $24,25$ -dihydroxylumisterol₃,

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24,25-dihydroxylumisterol₂, 1,24-dihydroxylumisterol₃,
1,24-dihydroxylumisterol₂, and 1,24-dihydroxy-25,26-
dehydrolumisterol₃.

5 34. The composition of claim 34, wherein said
carrier is effective for topical administration.

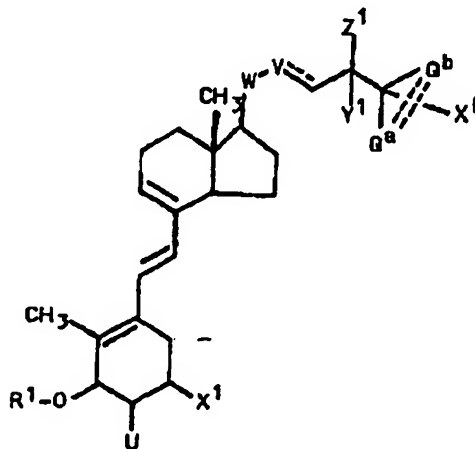
35. The composition of claim 32, further
comprising one or more sun screen agents.

36. The composition of claim 32, wherein said
carrier is effective for parenteral administration.

10 37. The composition of claim 32, wherein said
compound is present in an amount of 0.00001 to 10% by
weight.

15 38. The composition of claim 32, wherein said
compound is present in an amount of 0.0001 to 0.01% by
weight.

39. A composition comprising a pharmaceutically
acceptable carrier and a compound having the formula:



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wherein the bond between carbons C-22 and C-23 is single or double bond;

Y^1 is hydrogen, F, CH_3 , CH_2CH_3 or X^1 ;

U is hydrogen, -OH or -O-(C_2 - C_4 alkyl)-OH;

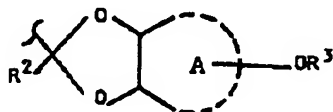
Z^1 is F, H or X^1 ;

Q^a is CF_3 or CH_2X^1 ;

Q^b is CF_3 or CH_3 ;

wherein X^1 is selected from the group consisting of hydrogen, -OH and OR^1 ;

wherein R^1 is hydrogen or a straight or branched chain glycosidic residue containing 1-20 glycosidic units per residue, or R^1 is an orthoester glycoside moiety of the Formula (III):



wherein A represents a glucofuranosyl or glucopyranosyl ring;

R^2 is hydrogen, lower alkyl, aralkyl, or aryl, with the proviso that aryl is phenyl or phenyl substituted by chloro, fluoro, bromo, iodo, lower C_1 - C_4 alkyl, C_1 - C_4 alkoxy; or naphthyl; and

R^3 is hydrogen or a straight or branched chain glycosidic residue containing 1-20 glycosidic units per residue;

W is $CH-CH_3$ or O; -

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V is CH₂ or O;

with the proviso that both W and V are not both O; and

"—" is either a single bond between Q^a and Q^b or a hydrogen atom on Q^a and Q^b;

5 wherein said compound is present in an amount effective to provide a vitamin D analog when said composition is administered to an individual.

10 40. The composition of claim 39, wherein said compound is 1alpha,25-dihydroxytachysterol₃, 1alpha,25-dihydroxytachysterol₂, 1alpha-hydroxytachysterol₃, 1alpha-hydroxytachysterol₂, 24,25-dihydroxytachysterol₃, 24,25-dihydroxytachysterol₂, 1,24-dihydroxytachysterol₃, 1,24-dihydroxytachysterol₂, and 1,24-dihydroxy-25,26-dehydrotachysterol₃.

15 41. The composition of claim 39, wherein said carrier is effective for topical administration.

42. The composition of claim 41, further comprising one or more sun screen agents.

20 43. The composition of claim 39, wherein said carrier is effective for parenteral administration.

44. The composition of claim 39, wherein said compound is present in an amount of 0.00001 to 10% by weight.

25 45. The composition of claim 39, wherein said compound is present in an amount of 0.0001 to 0.01% by weight.

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46. A method for providing a vitamin D analog to an individual which comprises administering to said individual the composition of claim 32 or 39 and exposing said individual to UV radiation.

5 47. The method of claim 46, wherein said composition is administered by topical means.

48. The method of claim 47, wherein said composition further comprises one or more sun screen agents.

10 49. The method of claim 46, wherein said composition is administered by intravenous means.

15 50. The method of claim 46, wherein said UV radiation is provided by sunlight of insufficient intensity and wavelength to effect the conversion of the corresponding provitamin D analog to the vitamin D analog.

51. The method of claim 46, wherein said UV radiation has a wavelength above 315 nm.

20 52. A method for treating decubitus or diabetic foot ulcers; ulcerative keratitis; psoriasis; wounds; inhibiting scar formation; glucocorticoid-induced decrease in calcium absorption; osteoporosis; senile decrease in calcium absorption; hypoparathyroidism; milk fever disease; turkey weak leg disease or
25 treating or preventing osteodystrophy due to an acquired or inherited defect in the metabolism of

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vitamin D; which comprises administering to said individual the composition of claim 32 or 39 and exposing said individual to UV radiation.

53. The method of claim 52, wherein said composition is administered by topical means.

54. The method of claim 53, wherein said composition further comprises one or more sun screen agents.

55. The method of claim 52, wherein said composition is administered by intravenous means.

56. The method of claim 52, wherein said UV radiation is provided by sunlight of insufficient intensity and wave-length to effect the conversion of the corresponding provitamin D analog to the vitamin D analog.

57. The method of claim 56, wherein said UV radiation has a wavelength above 315 nm.

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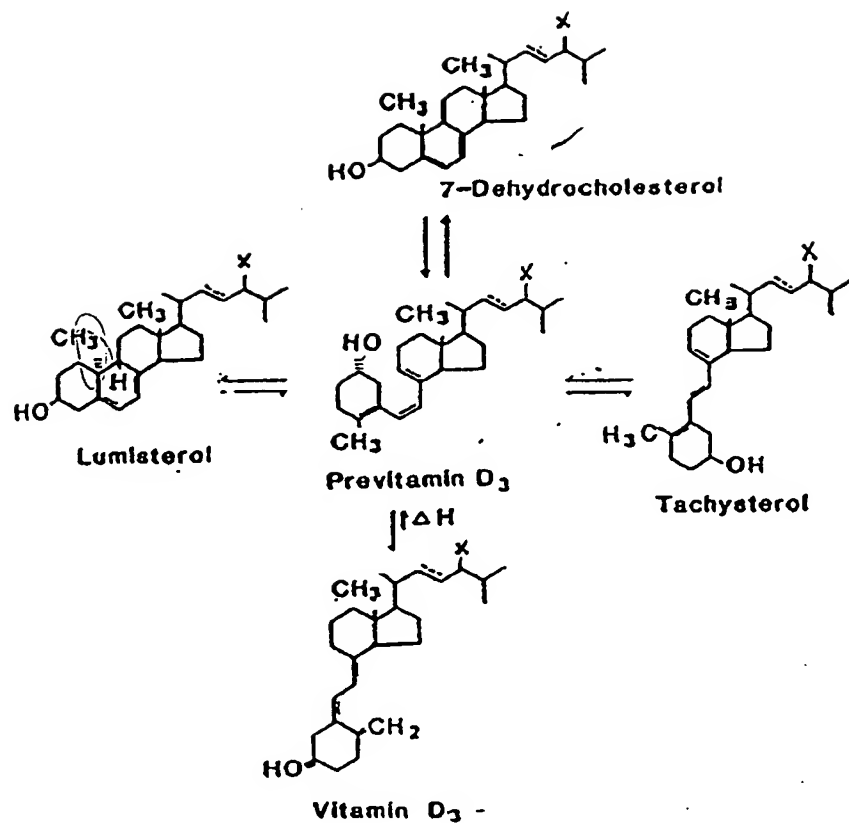
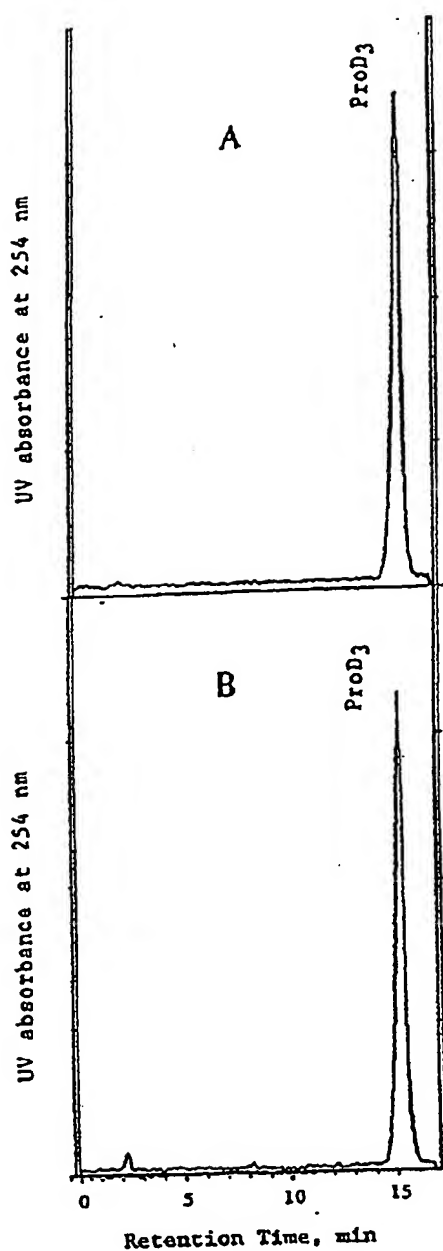
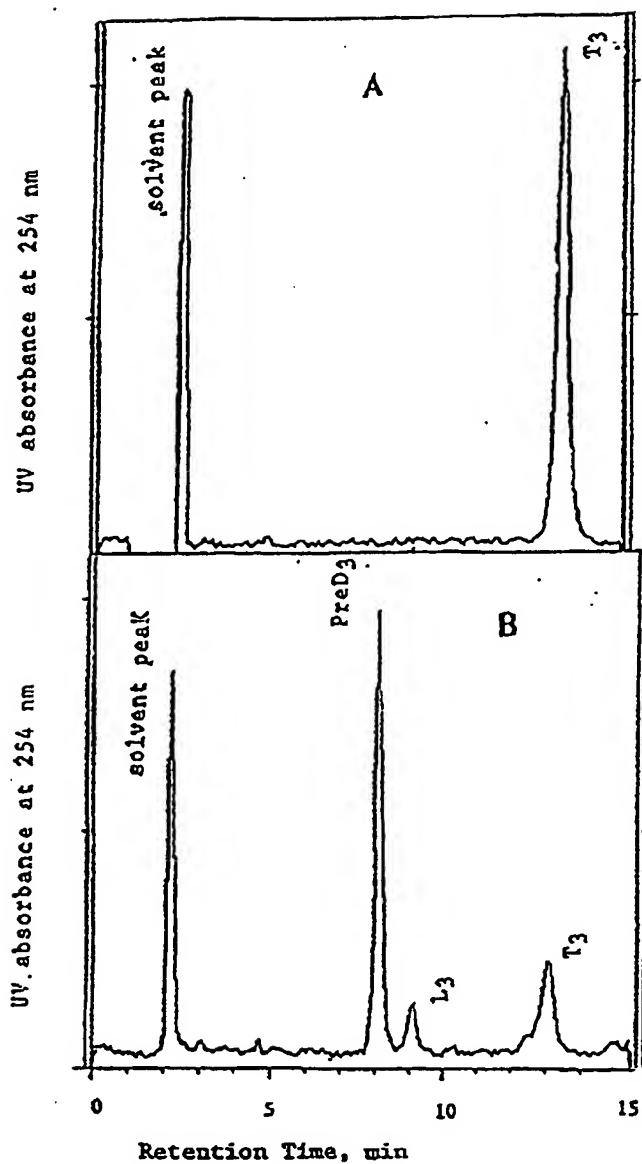


FIGURE 1



Exposure of ProvitaminD₃(ProD₃)
(10 µg/ml) to Sunlight on Nov.11,
1989 between 9:00-10:00 AM
A - Control. B - Exposed

FIGURE 2



Exposure of Tachysterol₃(T₃)(10 µg/ml) to Sunlight on
Nov.11,1989 between 9:00-10:00 AM

A - Control, B - Exposed

PreD₃ - Previtamin D₃, T₃- Tachysterol₃

FIGURE 3

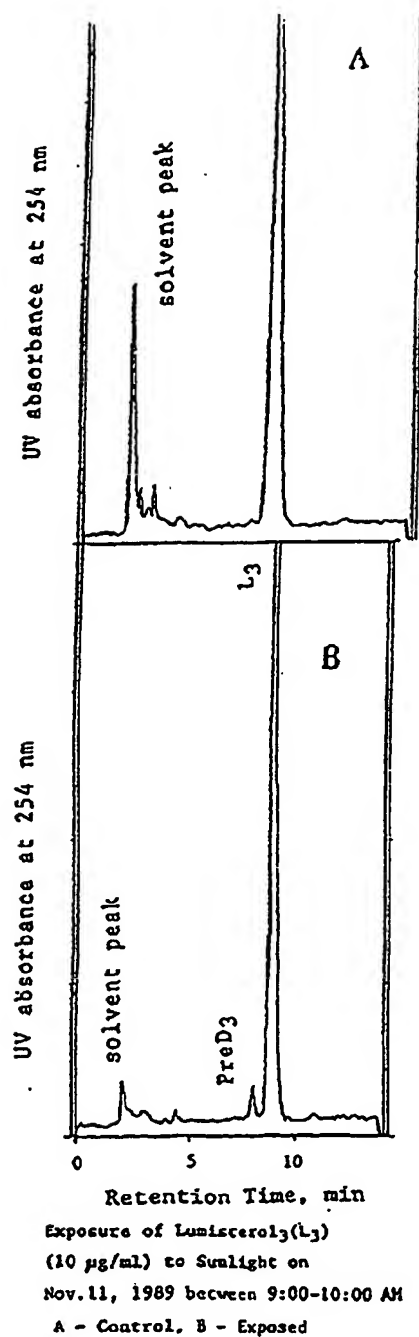


FIGURE 4

INTERNATIONAL SEARCH REPORT

International Application No. PCT/US91/04436

I. CLASSIFICATION & SUBJECT MATTER <small>See general classification in PCT/US91/04436, page 10</small>		
According to International Patent Classification (IPC) or to National Classification and IPC IPC(5): A61K 7/42, 44 31/59, 70, 715 US CL : 424/59, 60 514, 25, 54, 167, 171		
II. FIELDS SEARCHED		
<small>Minimum Documentation Searched</small> U.S. 424/59, 60 514/25, 54, 167, 171		
<small>Documentation Searched other than Minimum Documentation to the extent that such Documents are included in the Fields Searched</small> APS, CAS Online, Derwent: Compounds of the claims with terms such as SUNSCREEN, VITAMIN D, VITAMIN D DEFICIENCY, OSTEOMALACIA, OSTEOPOROSIS, CALCIUM, HYPOPARATHYROIDISM, WOUND, ULCER, DIABETES		
III. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of Document, with indication, where appropriate, of the relevant passages	Relevant to Claim No. *
Y	US, A, 3,702,810 (DE LUCA ET AL.) 14 NOVEMBER 1972 See column 1, lines 15-23.	8-14, 39-45
A	US, A, 4,230,701 (HOLICK ET AL.) 28 OCTOBER 1980 See entire document.	1-57
Y	US, A, 4,310,511 (HOLICK) 12 JANUARY 1982 See column 10, lines 10-67.	1-7, 20-38, 46-57
Y	US, A, 4,335,120 (HOLICK ET AL.) 15 JUNE 1982 See column 2, line 40-column 4, line 49; column 6, lines 33-56.	1-57
<p>* Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"A" document member of the same patent family</p>		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search 28 OCTOBER 1991		Date of Mailing of this International Search Report 07 NOV 1991
International Searching Authority ISA/US		Signature of Authorized Officer NGUYEN NGOC HO INTERNATIONAL DIVISION KIMBERLY JORDAN

Form PCT/ISA/E10 (second sheet) (Rev.11-87)